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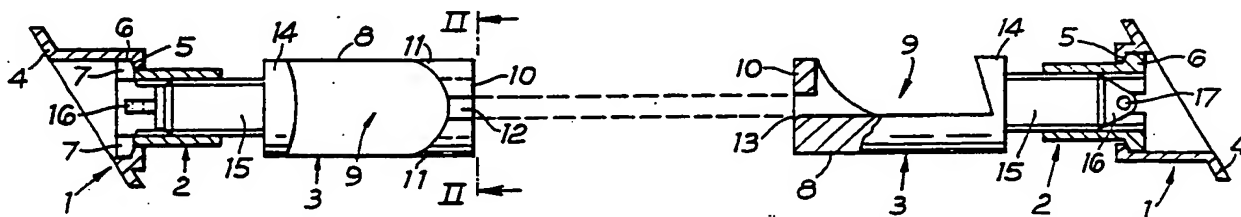
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<p>(21) International Application Number: PCT/GB91/01295 (22) International Filing Date: 30 July 1991 (30.07.91) (30) Priority data: 9016761.0 31 July 1990 (31.07.90) GB (71)(72) Applicant and Inventor: JOHNSON, David, Paul [GB/GB]; Woodland Lodge, 69 Cleveland Road, Failand, Bristol BS8 3UL (GB). (74) Agents: LAINÉ, Simon, James et al.; Wynne-Jones, Laine & James, 22 Rodney Road, Cheltenham, Gloucestershire GL50 1JJ (GB).</p>		<p>(81) Designated States: AT (European patent), AU, BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, KR, LU (European patent), NL (European patent), SE (European patent), US. Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>

(54) Title: IMPROVEMENTS RELATING TO SURGICAL DEVICES



(57) Abstract

A screw tensioning device is proposed for holding at least one end of a ligament graft. The device has a thimble (1) which mates in the mouth of a drilling through bone, a nut (2) captively seated in the thimble, and an anchorage element (3) with a screw threaded stud (12) which can engage the nut. Different forms of element are provided for different grafts, but each is adapted securely to hold one end. The anchorage element (3) with ligament attached is drawn through the drilling from the opposite side to the thimble (1) until the stud (12) engages the nut (2). That is turned by a tool (19) until the required tension is achieved.

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⁺ It is not yet known for which States of the former Soviet Union any designation of the Soviet Union has effect.

"Improvements relating to Surgical Devices"

This invention relates to surgical devices and is concerned with an adjustable tensioning assembly for ligaments.

5 The recent treatment of chronic knee instability due to rupture of the anterior cruciate ligament has been a mixture of ligament repair, augmentation or substitution by tendon transfer. The mediocre results and the inability of most athletes to return to their previous sporting activities in
10 full capacity has prompted intense research into prosthetic cruciate replacement (Johnson RJ et al. 1984). Initial research centred upon the use of the patella tendon or tensor fascia larter. Subsequent research centred upon carbon fibre or dacron ligaments as scaffolds for the slow
15 invasion by fibrous tissue (Butler DL et al 1985). Recently great interest has been shown in expanded polytetrafluoroethylene or "Gore-tex" ligaments. This ligament was first used in a multi centre research project in 1984 and the initial good results have resulted in its wider use. However, the
20 long term results¹ of independent assessment suggest an incidence of chronic knee effusion and late rupture of the Gore-tex graft. As a result, the most common technique at the present time is the use of an autogenous patella tendon graft with bone harvested from the patella and the tibial
25 tuberosity. The patella tendon graft and bone plugs are fed into tunnels within the femur and tibia and secured at each end.

One of the outstanding problems and the subject of much

discussion is the isometric placement of these ligaments. Theoretically isometric placement is possible where the ligament remains at the same tension throughout the range of knee flexion. However, this is difficult to achieve in practice. Many recent projects have defined the isometric points for insertion of the drill and have demonstrated the effect of incorrect placement. However, an unsolved problem is the fixation of the ends of the ligament with sufficient strength for early motion to be allowed. Different techniques include screws, baffles, bone blocks, staples, washers, screws and more recently toggles (Amis AA 1988. Good et al 1988). These all provide support which may be unreliable and which usually require protection or cautious use for up to one year to allow adequate fibres or bony ingrowth to provide secure fixation. This delays the return to activity, particularly sports, which leads to more muscle wasting and stiffness.

The final problem, which has yet received little attention, is correct tensioning of the ligament. For the ligament to be functional it not only has to be inserted isometrically but it has to be correctly tensioned in order to allow a full range of motion. In particular, it needs to be tight enough to give stability rather than being a check rein loaded only at the extremes of motion. With some ligaments, such as the Leeds Keio ligament, maintenance of tension during insertion is technically difficult, and early reports suggest some degree of laxity is often present post-operatively. No method of insertion is believed to be

currently available such that the tension may be incrementally increased while the range of motion and stability is continuously examined.

The aim of this invention is to enable a ligament to
5 be inserted and firmly held in position in an easy and certain manner, while allowing the tension in the ligament to be adjusted to an optimum. It may even be adjustable at a later operation if necessary. It should also be usable for a prosthetic ligament graft as well as an autogenous patella
10 tendon graft.

According to the present invention there is provided a tensioning device for ligament grafts, the device comprising an anchorage element for receiving and holding one end of a
15 ligament when it is under tension and leading from one end of said element, a bone engaging thimble, and screw means located by the thimble and co-operating with said element at its other end for drawing said element and said thimble together.

To fit such a device the bone is drilled through and
20 the anchorage element, with one end of the ligament engaged with it, is inserted through the bore from one side. The thimble is placed to abut the opposite side of the bone, having a portion which locates it partially within the other end of the bore. The screw means are then engaged and as
25 they are tightened the ligament is tensioned and the thimble is drawn more firmly against the bone.

Usually, there will be a pair of such devices with the other end of the ligament anchored in a similar manner,

alth ough screw-adjustability at both ends may not always be required.

In the preferred form, the thimble is generally of hollow cylindrical form having an external flange at one end and an internal shoulder at the other end, the flange being for abutment of bone around the mouth of a drilling in which the remainder of the device is received, and the shoulder being for retention of the screw means. The flange will generally be at a slant with respect to the axis of the cylinder, preferably in the range of 45° - 60° and the screw means will have an external projection for engagement of said internal shoulder of the thimble. The screw means may have a detent in its end for turning by a tool entered through the thimble, and conveniently it takes the form of a nut engageable on a threaded portion of said anchorage element.

The anchorage element will preferably be of generally cylindrical form co-axial with the screw means to fit snugly in the bore through the bone, but having a lateral recess to accept an end of the ligament. In one form a slot leads from said recess to said one to guide the ligament into leading substantially co-axially from said element, the portion in which the slot is formed providing an abutment for a bone fragment at the end of a harvested tendon. In another form, the recess has an upstanding abutment within the envelope of said element for the retention of the end of a prosthetic anterior cruciate graft.

For a better understanding of the invention one

embodiment will now be described, by way of example, with reference to the accompanying drawing, in which:

Figure 1 is a side view, partially in longitudinal section, of a ligament tensioning device,

5 Figure 2 is a section on the line II-II of Figure 1,

Figure 3 shows a lead wire for use with the device,

Figure 4 is a side view of a tensioning key for use with the device, and

10 Figure 5 shows views of an alternative carrier that could be incorporated in the device.

Reference will be made to inner and outer ends, these being with reference to the mid-point of the assembly.

At each end the tensioning device comprises a thimble 1 outermost, an intermediate nut 2, and a carrier 3 innermost. At the left hand end the carrier 3 is shown rotated through 90° from identical orientation with the carrier at the right hand end.

20 Each thimble 1 is of squat cylindrical form with its outer end angled and having an external flange 4. This angle is preferably between 45° and 60° to the axis. Internally, the thimble has a thickened portion towards its inner end to form a shoulder 5.

Each nut 2 is basically a hollow internally screw-threaded cylinder with an external rim 6 at its outer end, 25 the underside of which co-operates with the shoulder 5 of the associated thimble. At this end, diametrically opposite zones are cut away to form slots 7. At the inner end, there is an internal bevel to facilitate coupling to the carrier

3. The nuts 2 may be provided in various lengths.

Each carrier 3 has a main body 8 having a cylindrical envelope, but this has a middle portion cut away to form a recess 9. At its inner end, the wall 10 defining the limit
5 of the recess is shaped with re-inforcing shoulders 11 and is cut away in a T-shaped slot 12 as best seen in Figure 2. The cross arm of the T opens to the base of the recess 9 and is virtually diametral of the inner end of the carrier. Where it emerges to the end face, there is a rounded or
10 bevelled edge 13 to reduce chafe. At the outer end of the recess 9, the end wall 14 is undercut to contain the bone fragment. Beyond that there is a co-axial stud 15 externally screw threaded to mate with the nut 2. At its coned tip, which eases entry into the nut 2, the stud 12 has a trans-
15 verse lug 16 with a small hole 17.

For fitting this device the bone is drilled through with a bore corresponding to the envelope diameter of the carriers 3, which is the same as that of the non-flanged parts of the thimbles 1. A lead wire 18 is connected to one
20 of the carriers 3 using the hole 17. A suitable configuration for the wire 15 is shown in Figure 3. The long loop is then passed through the bore, and also through the nut 2 and thimble 1 which are lightly held in place at the other end of the bore. It can also be passed through a tensioning
25 key 19 (Figure 4) which is of hollow cylindrical form with a knurled portion 20 for a good grip, and two lugs 21 at one end to co-operate with the slots 7. The carrier 3, with a bone fragment located in the recess 9 and the attached

tendon 1 ading through the slot 12, is drawn up the bore until the stud 15 engages the nut 2, which can then be turned using the key 19 to complete the tensioning. Once that is done, the key can be removed, and then the lead wire
5 18.

This assumes that the ligament is already anchored at the other end. If it is not, then of course there will be no tension and the nut is simply done up a number of turns to ensure a good grip, while allowing for later adjustment.

10 The alternative carrier of Figure 5 differs by having a bollard 22 upstanding in the recess 9 and inclining towards the stud end. The undercut at that end is not required and both end walls have generally the same configuration with the T-slot opened out into a square one 23.
15 This bollard enables a prosthetic anterior cruciate graft to be coupled as an alternative to an autogenous patella tendon.

CLAIMS

1. A tensioning device for ligament grafts, the device comprising an anchorage element (3) for receiving and holding one end of a ligament when it is under tension and
5 leading from one end of said element, a bone-engaging thimble (1), and screw means (2) located by the thimble and co-operating with said element at its other end for drawing said element and said thimble together.

2. A device as claimed in Claim 1, characterised in
10 that the thimble (1) is generally of hollow cylindrical form having an external flange (4) at one end and an internal shoulder (5) at the other end, the flange (4) being for abutment of bone around the mouth of a drilling in which the remainder of the device is received, and the shoulder (5)
15 being for retention of the screw means (2).

3. A device as claimed in Claim 2, characterised in that the flange (4) is at a slant with respect to the axis of the thimble.

4. A device as claimed in Claim 2 or 3, characterised
20 in that the screw means (2) has an external projection (6) for engagement of said internal shoulder (4) of the thimble.

5. A device as claimed in Claim 2, 3 or 4, characterised in that the screw means (2) has a detent (7) in its end for turning by a tool (19) entered through the thimble (1).

25 6. A device as claimed in any preceding Claim, wherein the screw means (2) is a nut engageable on a threaded portion (15) of said anchorage element (3).

7. A device as claimed in any preceding Claim,

characterised in that the anchorage element (3) is of generally cylindrical form co-axial with the screw means (2) and has a lateral recess (9) to accept an end of the ligament.

5 8. A device as claimed in Claim 7, characterised in that a slot (12) leads from said recess (9) to said one end to guide the ligament into leading substantially co-axially from said element, the portion in which the slot is formed providing an abutment (10) for a bone fragment at the end of
10 a harvested tendon.

 9. A device as claimed in Claim 7, characterised in that the recess (9) has an upstanding abutment (22) within the envelope of said element for the retention of the end of a prosthetic anterior cruciate graft.

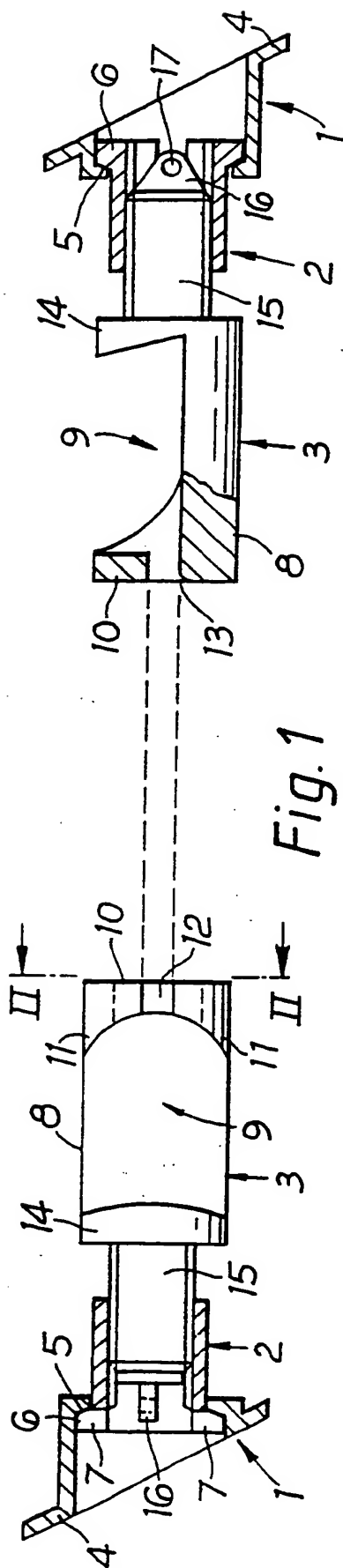


Fig. 1

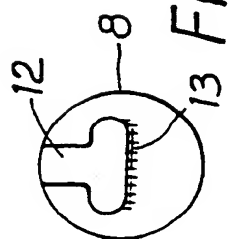


Fig. 2

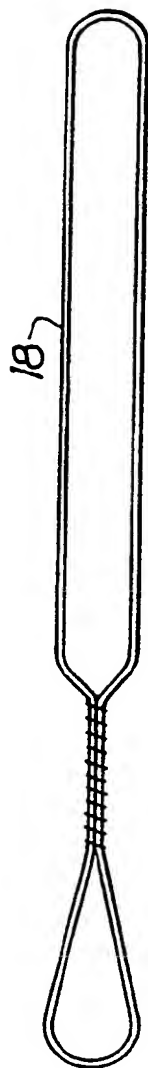


Fig. 3

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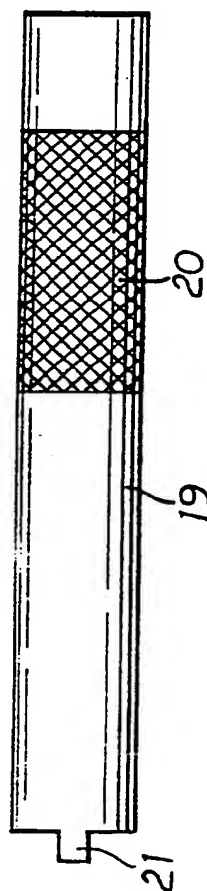


Fig. 4

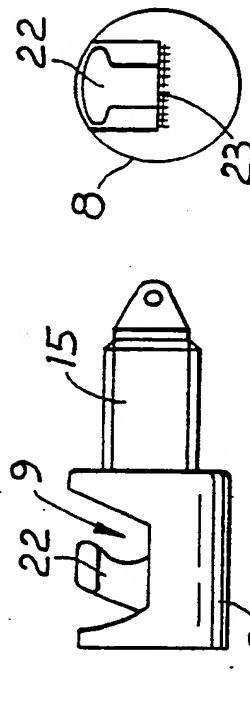


Fig. 5

INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 91/01295

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC Int.Cl. 5 A61F2/08		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
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Int.Cl. 5	A61F	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
X	FR,A,2 483 772 (RAMBERT) 11 December 1981	1,6
Y	see page 5, line 15 - line 19; figure 3 ---	2-5,7,8
Y	EP,A,0 232 049 (PFIZER) 12 August 1987 see page 12, paragraph 2; figure 9 ---	2-5
X	CH,A,267 810 (RUTSCHI) 1 July 1950 see page 2, line 37 - line 46; figure 4 ---	1,7
Y	US,A,3 973 277 (SEMPLE) 10 August 1976 see abstract; figures 1,2 ---	7,8
A	US,A,4 772 286 (GOBLE) 20 September 1988 ---	
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Patent document cited in search report	Publication date	Patent family member(s)	Publication date
FR-A-2483772	11-12-81	None	
EP-A-0232049	12-08-87	US-A- 4708132	24-11-87
		AU-B- 583972	11-05-89
		AU-A- 6796687	13-08-87
		CA-A- 1277453	11-12-90
		DE-U- 8701123	02-04-87
		JP-A- 62179454	06-08-87
CH-A-267810		None	
US-A-3973277	10-08-76	GB-A- 1465744	02-03-77
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		AU-A- 1618288	16-11-89
		EP-A- 0342281	23-11-89
DE-U-8809612	15-09-88	None	

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